

Fig. 1

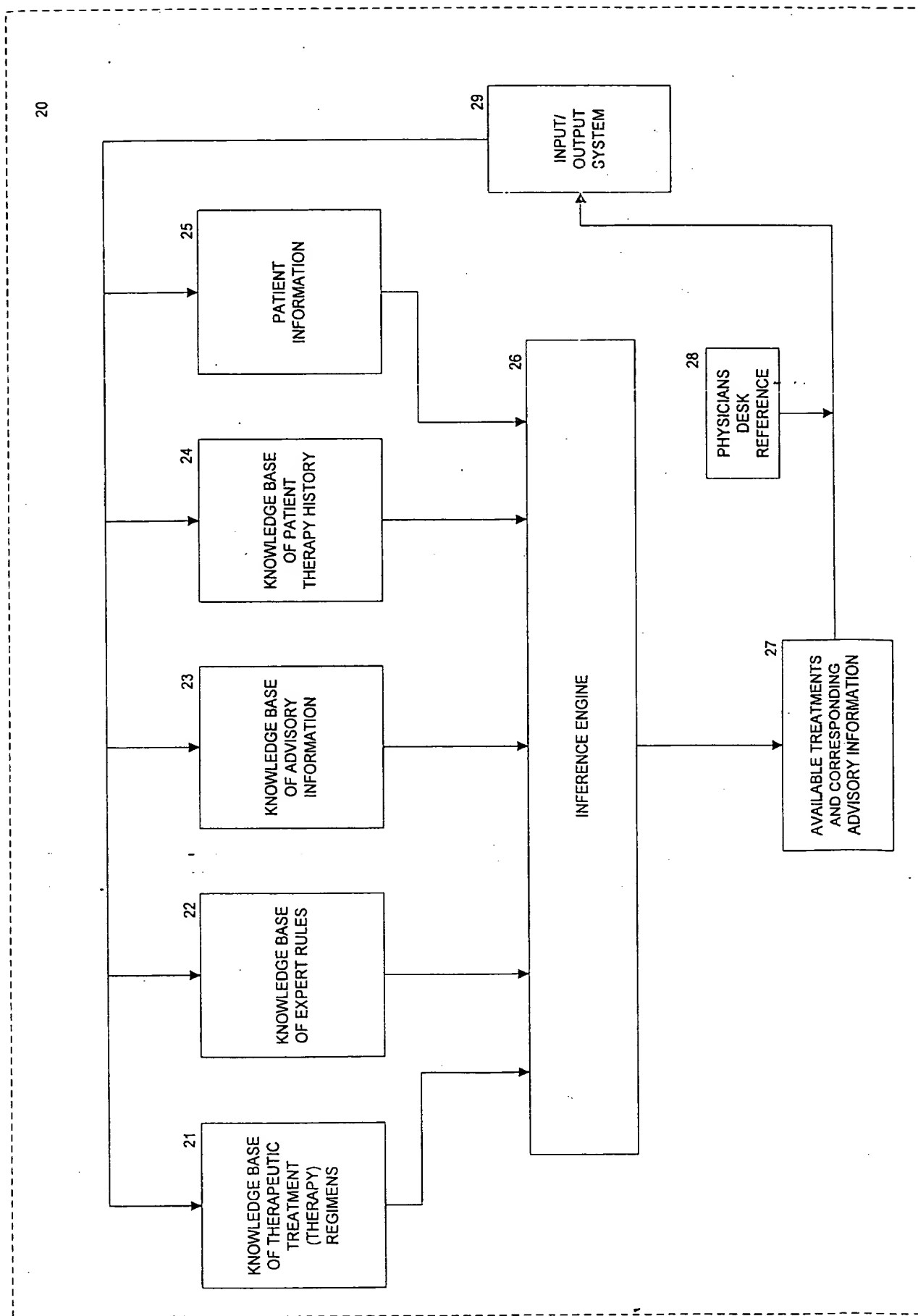
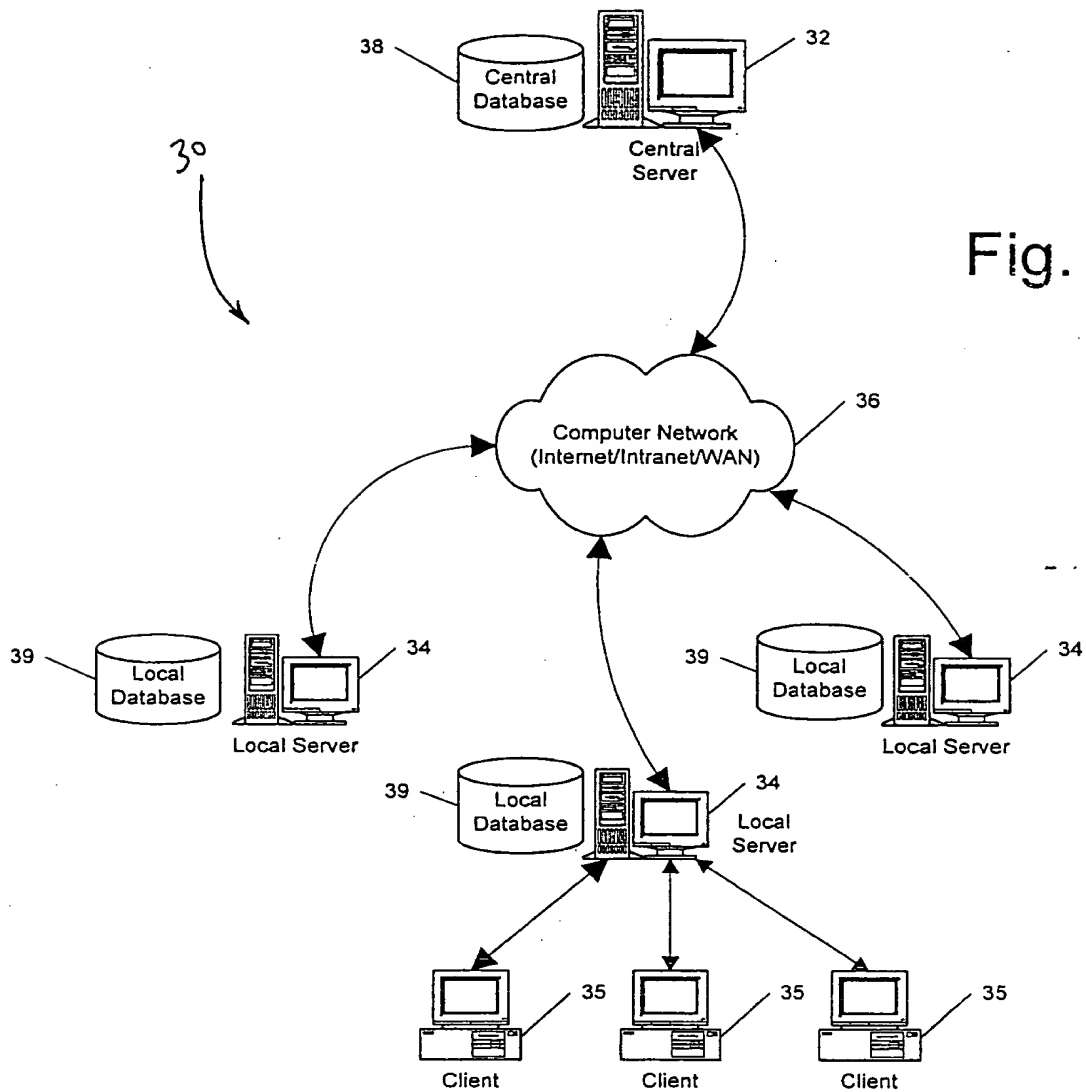


Fig. 2



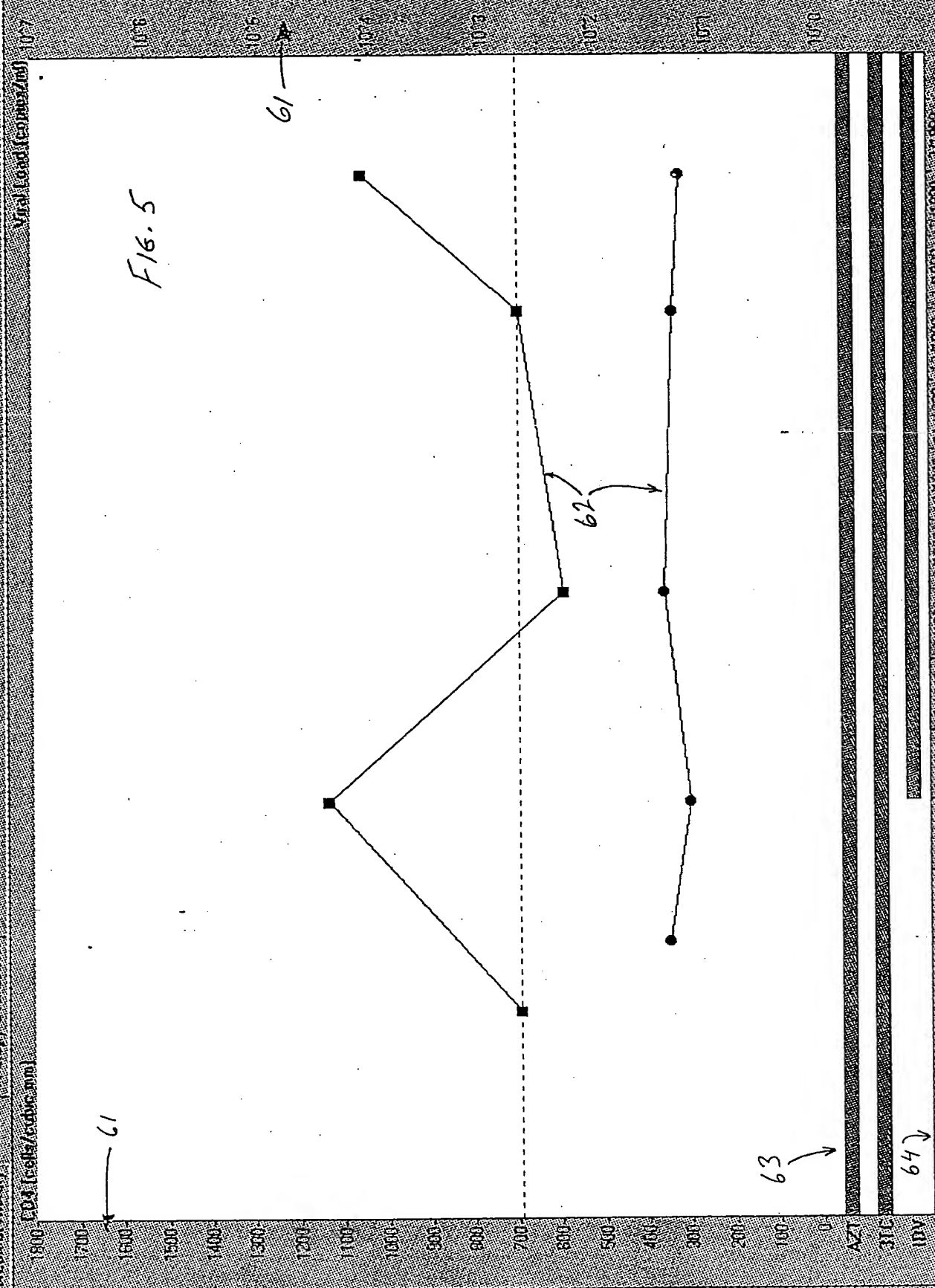


660

60a 70a

TPMS Patient

Medical History Chart Therapy Evaluation



TPMS

61 62 63 64

71 60a 72 73 70

**TPMS Patient**

Medical History: ☒ Chart ☒ Therapy Evaluation

Evaluate Current Therapy: ☒ AZT, 3TC, IDV ☐ Show 1 Drug Therapies ☒ Show 2 Drug Therapies ☐ Show 3 Drug Therapies ☐ Show Bested Therapies ☐ Show EAP Therapies

Therapy Options (10 of 17)

Therapy	El	Ad	Safety Considerations	Med	Dose	Freq	Cost
<input checked="" type="checkbox"/> AZT, d4T, NFV	2	2	ddl Renal dos. Adj. d4T Renal dos. Adj.	Y	q8h	15	\$30.38
<input type="checkbox"/> AZT, d4T, IDV	3	6	ddl Renal dos. Adj. d4T Renal dos. Adj. IDV Renal d...	Y	q8h	12	\$26.80
<input type="checkbox"/> AZT, d4T, RTV	4	7	ddl Renal dos. Adj. d4T Renal dos. Adj.	Y	q12h	18	\$34.06
<input type="checkbox"/> AZT, SQV-SGC, NFV	5	8	d4T Renal dos. Adj.	Y	q8h	29	\$45.60
<input checked="" type="checkbox"/> AZT, SQV-SGC, NFV	5	8	ddl Renal dos. Adj.	Y	q8h	31	\$42.24
<input type="checkbox"/> AZT, SQV-SGC, NFV	5	8	ddl Renal dos. Adj. tobramycin-ddC	Y	q8h	29	\$42.72
<input type="checkbox"/> AZT, d4T, NFV	8	8	ddl Renal dos. Adj. d4T Renal dos. Adj. tobramycin...	Y	q8h	13	\$30.86
<input type="checkbox"/> AZT, d4T, SQV-SGC	6	9	ddl Renal dos. Adj. d4T Renal dos. Adj.	Y	q8h	24	\$31.24

See More: ☐ Ther AB ☐ Top 10 ☒ Full Screen Evaluation

Antiretroviral Drug: ☒ AZT (Zalcitabine) ☒ ddI (Didanosine) ☒ ddC (Dideoxycytidine) ☒ 3TC (Lamivudine) ☒ d4T (Zalcitabine) ☒ ABC (Zalcitabine) ☒ IDV (Didanosine) ☒ SQV-HGC (Intravir/Indinavir)

Nucleoside Analogues (NRTI): ☒ AZT (Zalcitabine) ☒ ddI (Didanosine) ☒ ddC (Dideoxycytidine) ☒ 3TC (Lamivudine) ☒ d4T (Zalcitabine) ☒ ABC (Zalcitabine) ☒ IDV (Didanosine) ☒ SQV-HGC (Intravir/Indinavir)

Protease Inhibitors (PI): ☒ IDV (Didanosine) ☒ SQV-HGC (Intravir/Indinavir)

FIG-6

74

**Recommended Dosages**








- Vilex 125mg q12h (4 pills/day, \$4.22/day)
- Zerit 15mg q12h (2 pills/day, \$7.58/day)
- Crivian 800mg q8h (6 pills/day, \$15.00/day)

(\* indicates adjusted dosage)

**Warning - Resistance Notices**

- **d4T: Resistance Advisory:** Cross Resistance: The patient has at least one previous exposure to AZT that was greater than one year in duration. Previous AZT exposure can lessen the antiretroviral effect of d4T due to cross resistance. Therapies containing d4T have been ranked lower in their Adjusted Score by +3.
- **Resistance Advisory:** IDV: According to the last genotype data entered, the patient's virus currently has the following secondary mutation(s), (L101[P], I54V [P], and 184V [P]) which is/are associated with resistance to IDV. These mutations alone are not enough to preclude the use of IDV but they do indicate a trend in this direction. IDV is still an option but ongoing IDV use may result in a more rapid emergence of complete resistance. The Adjusted Score of IDV has been lowered by +3.

FIG. 7

Icon	Meaning
	Indicates that there were no critical alerts for the therapy, however, general warnings and advisories should be read in the Therapy Details box.
	Indicates that there were no critical alerts for the therapy, however, general warnings and advisories should be read in the Therapy Details box. The book indicates that therapy has been studied and a reference is available to review.
	Indicates a yellow alert. There is important information about this therapy that must be reviewed.
	Indicates a yellow alert. There is important information about this therapy that must be reviewed. The book indicates that therapy has been studied and a reference is available to review.
	Indicates a red alert, which means critical and possible life-threatening situation may exist or may be created with this therapy. Information in the Therapy Details section must be read for this therapy to be considered.
	Indicates a red alert, which means critical and possible life-threatening situation may exist or may be created with this therapy. Information in the Therapy Details section must be read for this therapy to be considered. The book indicates that therapy has been studied and a reference is available to review.
	Indicates the therapy is not recommended.

F16-8

73a 78 73b

WASP Patient

Medical History | Chart | Therapy Evaluation

Therapy Being Evaluated

AZT, ddI, SQV, RTV

Use Current Therapy

Stop Therapy

**STOP! DRUG INTERACTION RED ALERT IS STOP!**

Read the following Red Drug Contra-Indication Alerts for this therapy:

Drug Interaction Alert

• Patient is currently taking citalopram, co-administration of Norvir (Ritonavir/RTV) with certain non-sedating antihistamines, sedative hypnotics or antiarrhythmics may result in potentially serious and/or life-threatening adverse events due to possible effects of Norvir (Ritonavir/RTV) on the hepatic metabolism of certain drugs. If Norvir (Ritonavir/RTV) can produce large increases in plasma concentrations of certain highly metabolized drugs, Norvir (Ritonavir/RTV) should not be administered with alprazolam, amiodarone, astemizole, bupropion, disipride, dioxepane, dioxepine, diazepam, encainide, etizolam, flunitrazepam, flurazepam, meprobamate, midazolam, piroxicam, propofol, propofol, quinine, rifabutin, rifampin, terfenadine, triazolam or zolpidem. Patient is taking citalopram and in order to use this therapy, that drug should be replaced with a non-contraindicated substitute. Consult your physician.

Dosages

- Ritonavir 300mg q12h (2 pills/day, \$9.56/day)
- Zidovudine 125mg q12h (4 pills/day, \$4.22/day)
- Zalcitabine 400mg q12h, taken within 2 hours after a full meal (4 pills/day, \$1.47/day)
- Nevirapine 400mg q12h (8 pills/day, \$1.44/day)

Dosage Adjustments

The following dosage adjustments messages apply to this therapy:

• Dosage Notice: This therapy contains both zalcitabine and zidovudine. When zalcitabine and zidovudine are used together the dosage of each drug is reduced by 1/2. The dosage for these drugs has been set accordingly. Do not use. Consult your physician.

Warnings and Advisories

The following Warnings and Advisories apply to Invirase (zalcitabine/SQV):

• Drug Interaction Information: Compounds that are substrates of CYP3A4 (e.g., calcium channel blockers, clindamycin, dapsone, quinidine, in zalcitabine) may have elevated plasma concentrations when administered with Invirase (zalcitabine/SQV); therefore, patient should be monitored for toxicity associated with such drugs when taking Invirase (zalcitabine/SQV). Consult your physician.

73

73d

73e

73f

73g



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76

Therapy Options

Therapy	Eff	Ad	Safety
1 d4T, 3TC, IDV	1	1	
1 AZT, 3TC, IDV	1	1	
2 d4T, 3TC, NFV	1	1	
1 AZT, 3TC, NFV	1	1	
2 d4T, 3TC, NFV			
1 AZT, 3TC, NFV			
1 ddI, d4T, 3TC			
2 d4T, 3TC, NFV			
2 d4T, 3TC, NFV			

Therapy B

Evaluated

General

- Vi
- Me

Show Abstract for Retrovir

Show Abstract for Efavir

Show Abstract for Viracept

Show Therapy Study

Print Details for AZT, 3TC, NFV

Print Top 10 Therapy Option Details

Hide Column "Eff"

Hide Column "Ad"

Hide Column "Safety Considerations"

Show Column "Med"

Show Column "Drug"

Hide Column "Freq"

Hide Column "Pills"

Hide Column "Cost"

90

FIG. 9

General		Birth Date		Gender		TEWS Number		Entry		Comment/Propose		Date		Value	
Patient Id: demo1		1/1/1960		Male				<input type="checkbox"/> Entry <input checked="" type="checkbox"/> Comment/Propose				3/3/1999		55.00	
Physician:								<input type="checkbox"/> Entry <input checked="" type="checkbox"/> Comment/Propose				3/3/1999		55.00	
Physician:								<input type="checkbox"/> Entry <input checked="" type="checkbox"/> Comment/Propose				3/3/1999		55.00	
Physician:								<input type="checkbox"/> Entry <input checked="" type="checkbox"/> Comment/Propose				3/3/1999		55.00	
Physician:								<input type="checkbox"/> Entry <input checked="" type="checkbox"/> Comment/Propose				3/3/1999		55.00	
Physician:								<input type="checkbox"/> Entry <input checked="" type="checkbox"/> Comment/Propose				3/3/1999		55.00	
Physician:								<input type="checkbox"/> Entry <input checked="" type="checkbox"/> Comment/Propose				3/3/1999		55.00	
Physician:								<input type="checkbox"/> Entry <input checked="" type="checkbox"/> Comment/Propose				3/3/1999		55.00	
Physician:								<input type="checkbox"/> Entry <input checked="" type="checkbox"/> Comment/Propose				3/3/1999		55.00	
Physician:								<input type="checkbox"/> Entry <input checked="" type="checkbox"/> Comment/Propose				3/3/1999		55.00	
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Physician:								<input type="checkbox"/> Entry <input checked="" type="checkbox"/> Comment/Propose				3/3/1999		55.00	
Physician:								<input type="checkbox"/> Entry <input checked="" type="checkbox"/> Comment/Propose				3/3/199			

FIG. 10A

W



60

60a

70a

TPMS Patient

Medical History Chart Therapy Evaluation

Viral Load (copies/mL)

1800-1700-1600-1500-1400-1300-1200-1100-1000-900-800-700-600-500-400-300-200-100-0

61

FIG. 10C

61

62

63

64

12/1997 1/1998 2/1998 3/1998 4/1998 5/1998 6/1998 7/1998 8/1998 9/1998 10/1998 11/1998 12/1998 1/1999 2/1999 3/1999

TPMS +1

Navigation icons: Home, Back, Forward, Stop, Print, Web, Mail, etc.





50

Fig. 11A

TPMS Patient 60a 70a

Medical History Chart Therapy Evaluation

<b>General</b>		<b>Weight (kg)</b>		<b>Date</b>		<b>Value</b>	
Patient (d) [ARV naive]		1/5/1998		2/1/1999		73.00	
<b>Physician</b>		<b>Gender</b>		<b>Weight (kg)</b>		<b>Date</b>	
[ ]		Male		2/1/1999		73.00	
<b>TPMS Number</b>		<b>Print</b>		<b>Save</b>		<b>Yes</b>	
[ ]		[ ]		[ ]		[ ]	
<b>CD4 and Viral Load</b>							
<b>CD4</b>		<b>Specimen Date</b>		<b>Value</b>		<b>Prev Value</b>	
[ ]		2/20/1999		350		375	
<b>Current Viral Load</b>		<b>Specimen Date</b>		<b>Value</b>		<b>Prev Value</b>	
[ ]		2/20/1999		31000		[ ]	
<b>Previous Viral Load</b>		<b>Specimen Date</b>		<b>Value</b>		<b>Prev Value</b>	
[ ]		12/29/1998		19000		[ ]	
<b>HIV Genotype</b>							
<b>Phenotype</b>		<b>Specimen Date</b>		<b>Value</b>		<b>Prev Value</b>	
[ ]		[ ]		[ ]		[ ]	
<b>Allergy/Hyper</b>		<b>Specimen Date</b>		<b>Value</b>		<b>Prev Value</b>	
[ ]		[ ]		[ ]		[ ]	
<b>Intolerance</b>		<b>Specimen Date</b>		<b>Value</b>		<b>Prev Value</b>	
[ ]		[ ]		[ ]		[ ]	
<b>Hemoglobin</b>							
<b>Specimen Date</b>		<b>Value (g/dl)</b>		<b>Date</b>		<b>Value</b>	
[ ]		2/1/1999		12.50		[ ]	
<b>Neutrophils</b>							
<b>Specimen Date</b>		<b>Value (cells/cubic mm)</b>		<b>Date</b>		<b>Value</b>	
[ ]		2/1/1999		1350		[ ]	
<b>Hepatic Function</b>							
<b>Specimen Date</b>		<b>AST/SGOT (U/L)</b>		<b>ALT/SGPT (U/L)</b>		<b>Renal Function</b>	
2/1/1999		35		35		[ ]	
<b>Specimen Date</b>		<b>Diastolic</b>		<b>Systolic</b>		<b>Est Creatinine</b>	
2/1/1999		No		1.00		110.51	
<b>Current ARV Therapy</b>							
<b>Drug</b>		<b>Dose</b>		<b>Frequency</b>		<b>Start Date</b>	
Prozac Pulvules & Liquid, O...		oral		10/5/1998		[ ]	
Bactrim DS Tablets		oral		12/8/1998		[ ]	
<b>Non-ARV Drugs</b>							
<b>Drug</b>		<b>Dose</b>		<b>Frequency</b>		<b>Start Date</b>	
[ ]		[ ]		[ ]		[ ]	



70a

Medical History Chart Therapy Evaluation

General  
 Patient ID: ARV naive1  
 Birth Date: 1/5/1958  
 Gender: Male  
 Weight (kg): 73.00  
 Date: 2/1/1999  
 Solid Dosage: Yes  
 Comment Pop-up: ☐ Empty ☒ Comment Pop-up

CD4 and Viral Load  
 CD4 (cells/mm<sup>3</sup>): 276  
 Current Viral Load: 276  
 Previous Viral Load: 12  
 HIV Genotype: ☐ H ☐ H ☐ H  
 Phenotype: ☐ H ☐ H ☐ H  
 Allergy/Hyper: ☐ H ☐ H ☐ H  
 Intolerance: ☐ H ☐ H ☐ H  
 Hemoglobin: ☐ H ☐ H ☐ H  
 Specimen Date: 2/1/1999  
 Neutrophils: ☐ H ☐ H ☐ H  
 Specimen Date: 2/1/1999  
 Hepatic Function: ☐ H ☐ H ☐ H  
 Specimen Date: 2/1/1999  
 ALT: 35

Boundry and Prequalification Messages

Please be aware that the following hourly and prequalification conditions currently apply to this patient.

- **Therapy Initiation:** Current treatment guidelines recommend initiation of antiRetroviral therapy for HIV -infected patients with HIV RNA (viral load) concentrations greater than 20,000 copies/ml (10,000 E<sub>q</sub>/ml bDNA) or CD4 counts less than 300 cells/ $\mu$ L (Ann. Int. Med., 1998). PreQualM, Commentary61
- **Combination Therapy Recommended:** Experts agree that the goal of antiRetroviral therapy should be to reduce the viral load to as low a level as possible for as long as possible. Initiation of therapy with a combination containing 2 nucleoside reverse transcriptase inhibitors (NRTI's) and a potent protease inhibitor have been shown to provide enhanced clinical benefit versus 2 drug combinations with regard to reduction in viral load and improved clinical outcomes. PreQualM, Commentary66

MBZ

Fig. 113

TPMS



Therapy Being Evaluated: None

Therapy Dates (10 of 613)

Therapy	Elig	Ad	Safety Discontinuation	Reg	FN	Cost
<input checked="" type="checkbox"/> AZT, ddI, 3TC, SQV-SGC	1	1		q8h	26	\$43.46
<input checked="" type="checkbox"/> ddI, 3TC, NFV	1	1		q8h	13	\$34.78
<input checked="" type="checkbox"/> AZT, 3TC, IDV	1	1		q8h	10	\$32.24
<input checked="" type="checkbox"/> AZT, 3TC, NFV	1	1		q8h	13	\$35.81
<input checked="" type="checkbox"/> ddI, 3TC, IDV	1	1		q8h	10	\$31.20
<input checked="" type="checkbox"/> AZT, ddI, RTV, DLV	2	2	DLV+RTV	q8h	30	\$45.99
<input checked="" type="checkbox"/> ddI, ddI, IDV, NVP	2	2		q8h	17	\$42.55
<input checked="" type="checkbox"/> ddI, 3TC, RTV	2	2		q12h	16	\$38.46
<input checked="" type="checkbox"/> AZT, ddI, RTV, NVP	2	2		q12h	20	\$47.10

See More: See All [10] Full Screen Evaluation

Therapy Being Evaluated: None

- **WARNING::** Before initiating any antiRetroviral treatment regimen, the complete product information for each therapeutic component should be consulted.

CmtGenY, Commentary35

- **Viral Load Testing Required:** Viral load testing should be repeated 21-35 days after initiation of, or a change of, antiRetroviral therapy to evaluate therapeutic efficacy and patient compliance. CmtGenY, Commentary65

- **Therapy Initiation:** Current treatment guidelines recommend initiation of antiRetroviral therapy for HIV-infected patients with HIV RNA (viral load) concentrations greater than 20,000 copies/ml (10,000 E<sub>q</sub>/ml bDNA) or CD4 counts less than 500 cells/ $\mu$ l. (Ann. Int. Med., 1998). P<sub>re</sub>QualM, Commentary61

- **Combination Therapy Recommended:** Experts agree that the goal of antiRetroviral therapy should be to reduce the viral load to as low a level as possible for as long as possible. Initiation of therapy with a combination containing 2 nucleoside reverse transcriptase inhibitors (NRTI's) and a potent protease inhibitor have been shown to provide enhanced clinical benefit versus 2 drug combinations with regard to reduction in viral load and improved clinical outcomes. P<sub>re</sub>QualM, Commentary66

Show 3 Drug Therapies ☒ Show 3 Drug Therapies ☒ Show 4 Drug Therapies ☒ Show 4 Drug Therapies ☒

Antiretroviral Drug

Nucleoside Analogues (NRTI)

- ☐ AZT (Retrovir/zidovudine)
- ☐ ddI (Videx/ddanosine)
- ☐ ddC (Hivid/zalcitabine)
- ☐ 3TC (Epivir/lamivudine)
- ☐ d4T (Zerit/stavudine)
- ☐ ABC (Ziagen/abacavir)

Protease Inhibitors (PI)

- ☐ IDV (Crixivan/indinavir)
- ☐ SQV-HGC (Invirase/saquinavir)
- ☐ SQV-SGC (Famvirase/saquinavir)

Used as Parent Therapy

W1

A1

FIG-11C

A2

A3



Therapy Being Evaluated  
AZT, ddI, RTV, DLV

Use as Directed Therapy

Show Therapies

## Recommended Dosages

- Retrovir 300mg q12h (2 pills/day, \$9.56/day)
- Videx 200mg q12h (4 pills/day, \$6.78/day)
- Norvir 600mg q12h (12 pills/day, \$22.26/day)
- Rescriptor 400mg q8h (12 pills/day, \$7.39/day)

Fig. 11D

- AZT: Interrupt Retroviruse if anemia and/or neutropenia develops. More Info 036 DosGenA, Commentary36
- ddI: When treatment with other drugs known to cause pancreatic toxicity is required (for example, IV pentamidine), suspension of Videx should be considered. CmtGenA, Commentary13
- ddI: If patients develop symptoms of neuropathy, Videx therapy should be interrupted. DosGenB, Commentary40
- ddI: Clinical signs suggestive of pancreatitis should prompt dose suspension of Videx and careful evaluation of the possibility of pancreatitis. Only after pancreatitis has been ruled out should dosing be resumed. DosGenB, Commentary39
- DLV: Skin rash attributable to Rescriptor may occur during first 21 days. More Info 034 CmtGenS, Commentary54

- ddI: Videx should not be administered with a prescription antibiotic containing any form of tetracycline. CmtGenA, Commentary15
- ddI: Plasma concentrations of some quinolone antibiotics are decreased when administered with antacids containing magnesium or aluminum. Therefore, doses of quinolone antibiotics should not be administered within 2 hours of taking Videx. CmtGenA, Commentary16
- RTV: Monitor for decreased AUC of Norvir and associated adverse events when concomitant with use of drugs that increase CYP3A activity (including tobacco). More Info 026 CmtGenH, Commentary26

it regimen, the complete product information for each therapeutic component should be consulted.

- **WARNING:** Current treatment guidelines recommend initiation of antiRetroviral therapy for HIV-infected patients with HIV RNA (viral load) concentrations greater than 20,000 copies/mL (10,000 E<sub>q</sub>/mL bDNA) or CD4 counts less than 500 cells/ $\mu$ L (Ann. Int. Med., 1998). PreQualM, Commentary61
- **Viral Load Testing:** Current treatment guidelines recommend initiation of antiRetroviral therapy for HIV-infected patients with HIV RNA (viral load) concentrations greater than 20,000 copies/mL (10,000 E<sub>q</sub>/mL bDNA) or CD4 counts less than 500 cells/ $\mu$ L (Ann. Int. Med., 1998). PreQualM, Commentary61

and patient compliance. CmtGenY, Commentary65

- **Therapy Initiation:** Current treatment guidelines recommend initiation of antiRetroviral therapy for HIV-infected patients with HIV RNA (viral load) concentrations greater than 20,000 copies/mL (10,000 E<sub>q</sub>/mL bDNA) or CD4 counts less than 500 cells/ $\mu$ L (Ann. Int. Med., 1998). PreQualM, Commentary61

- **Combination Therapy Recommended:** Experts agree that the goal of antiRetroviral therapy should be to reduce the viral load to as low a level as possible for as long as possible. Initiation of therapy with a combination containing 2 nucleoside reverse transcriptase inhibitors (NRTIs) and a potent protease inhibitor have been shown to provide enhanced clinical benefit versus 2 drug combinations with regard to reduction in viral load and improved clinical outcomes. PreQualM, Commentary66

660

600 700

TPMS Patient

Medical History Chart Therapy Evaluation

CD4 (cells/cubic mm)

Viral Load (copies/ml)

10<sup>7</sup>

10<sup>6</sup>

10<sup>5</sup>

10<sup>4</sup>

10<sup>3</sup>

10<sup>2</sup>

10<sup>1</sup>

10<sup>0</sup>

10<sup>-1</sup>

10<sup>-2</sup>

10<sup>-3</sup>

10<sup>-4</sup>

10<sup>-5</sup>

10<sup>-6</sup>

10<sup>-7</sup>

10<sup>-8</sup>

10<sup>-9</sup>

10<sup>-10</sup>

10<sup>-11</sup>

10<sup>-12</sup>

10<sup>-13</sup>

10<sup>-14</sup>

10<sup>-15</sup>

F16-12A



A12

Phenotypic Resistance to 3TC  
from 3/15/1999 to present

3TC

d4T

NVP

AZT

ddV

ddC

ddI

ddE

ddF

ddG

ddH

ddI

ddJ

ddK

ddL

ddM

TPMS +1

100

100

100

100

100

100

100

100

100

100

100

100

100

100

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# TPMS Patient

Medical History | Chart | Therapy Evaluation

Evaluate Current Therapy > 3TC, d4T, NVP

Therapy	Eff	Ad	Safety Considerations	Freq	Pts	Cost
Δ d4T, d4T, NVP	2	2	Rifabutin+NVP	q8h	15	\$33.88
● d4T, d4T, EFV	5	5		q12h	9	\$28.44
Δ d4T, NVP, EFV	5	5	Rifabutin+NVP	q8h	16	\$38.50
Δ d4T, NVP, EFV	5	5	Rifabutin+NVP	q8h	14	\$40.24
Δ ddC, NVP, EFV	5	7	Rifabutin+NVP	q8h	15	\$38.77
● ddC, d4T, EFV	5	7		q8h	8	\$28.71

See More | See All | Top 10 | Full Screen Evaluation

Therapy Being Evaluated 3TC, d4T, NVP

## III THERAPY REJECTED III

*This therapy was rejected for the following reason(s) Additional information about the therapy is provided but this therapy is NOT advisable*

- Viramune (nevirapine/NVP) Resistance Advisory: According to the last genotype data entered, the patient's virus currently has mutation(s) which is/are associated with resistance to Viramune. FilMue, Rejection54
- Resistance Advisory: According to the last genotype data entered, the patient's virus currently has the following mutations; M184V [RT]. The genotype test displays evidence of the M184V/M184I mutation which is associated with resistance to 3TC. However, this mutant has increased sensitivity to the antiretroviral activity of AZT and ADV so an AZT/3TC or AZT/ADV combination is still useable. Therefore combinations which contain AZT/3TC and AZT/ADV are shown as therapy options although these therapies have been ranked down +5 in favor of three drug combinations with no resistant mutants. FilMueB, Rejection01
- Efavir and Viramune Resistance Advisory: The patient's last phenotypic assay demonstrates phenotypic resistance to Efavir and Viramune, therefore, therapies containing Efavir and Viramune are not recommended at this time. FilRes-C, Rejection#2

Fig. 12 B

CAUTION

YELLOW ALERT

CAUTION

W3

- NVPΔ: Drug Interaction Alert: Patient is currently taking rifabutin and there is insufficient data to assess whether dose adjustments are necessary. These drugs

TPMS



How To

FIG-12C

S4C

W3

5/1999  
5/1999

• **NPVA**: Drug Interaction Alert. Patient is currently taking nifedipine and there is insufficient data to assess whether dose adjustments are necessary. These drugs should only be used in combination if clearly indicated and with careful monitoring. CmtDIP, Commentary33

**Medical History** | **Chart** | **Therapy Evaluation**

**General**

Patient ID: Features1 Birth Date: 1/1/1960 TPMS Number: 1/28/1999 Date: 1/28/1999 Value: 60.00

Phyician: Patient Gender: Male Print Save

**CD4 and Viral Load**

CD4 Specimen Date: 3/15/1999 Value: 240 Specimen Date: 1/28/1999 Free Value: 265

Current Viral Load Specimen Date: 3/15/1999 Value: 21500 C/mL

Previous Viral Load Specimen Date: 1/28/1999 Value: 2800 C/mL

**HIV Genotype** Specimen Date: 3/15/1999 Value: L101(P), M46(P), I54V(P), V82A(P), M41L(P), Y181

**Neutrophils** Specimen Date: 1/28/1999 Value: 15.00

**Hemoglobin** Specimen Date: 1/28/1999 Value: 15.00

**Neutropathy** Specimen Date: 1/28/1999 Value: No

**Pancreatitis** Specimen Date: 1/28/1999 Value: No

**Hepatic Function** Specimen Date: 1/28/1999 Value: 25

**Renal Function** Specimen Date: 1/28/1999 Value: 25

**AIDS Diagnostic** Date: 1/28/1999 Value: Yes

**Current ARV Therapy** 1/28/1999 3TC, D4T, NVP

**Non-ARV Drugs**